UIA-T / UIA-T LEPHILE	HIV-1	/ HIV-2 Testing
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	HIV-1 / HIV-2 Testing
Toot	Ovalitative assert for the detection of 100/4 and 100/2 antihadias and
Test	Qualitative assay for the detection of HIV-1 and HIV-2 antibodies and
Description	HIV-1 p24 antigen in human serum or plasma
Test Use	Screening assay to aid in the diagnosis of infection with HIV-1 and/or HIV-2
Test	Virology
Department	Phone: (860) 920-6662, FAX (860) 920-6661
Methodology	Enzyme immunoassay (EIA) and HIV antibody differentiation assay
Availability	Daily, Monday-Friday
Specimen	1 mL serum
Requirements	1 mL plasma (acceptable anticoagulants include EDTA, sodium and lithium
	heparin, sodium citrate, CPD, CPDA-1 and ACD)
Collection	To obtain collection kit, refer to Collection Kit Ordering Information.
Kit/Container	
Collection	Standard venipuncture
Instructions	
Specimen	Store specimen at 2-8° C. Specimens must be received within 7 days of
Handling &	collection.
Transport	Transport with an ice pack coolant (preferred) or at ambient temperature.
	Avoid temperature extremes.
Unacceptable	Unlabeled specimens
Conditions	Specimens that have leaked or containers that have broken in transit
	Specimens received more than 7 days after collection
Requisition	Clinical test requisition (select HIV-1 / HIV-2 Testing)
Form	
	Name and address of submitter (and/or Horizon profile #)
Required	Patient name or identifier, town of residence (city, state, zip), date of birth
Information	Specimen type or source, date collected, test requested
	Please ensure patient name on requisition matches that on the specimen.
	A negative test result does not preclude the possibility of exposure to or
	infection with HIV-1 and/or HIV-2.
	The performance of this assay has not been established for children less
Limitations	than 2 years of age since maternal IgG frequently persists for as long as
	18 months after birth. Supplemental assays designed for neonatal
	specimens may be helpful in resolving such cases, including HIV nucleic
	acid tests or viral culture.
	Confirmatory testing is performed on repeatedly reactive specimens
	using an HIV antibody differentiation assay.
Additional	Specimens found to be positive by the submitter will be tested in
Comments	triplicate using the HIV-1 / HIV-2 Combo Antigen / Antibody assay.
	Confirmatory testing will only be performed on specimens found to be
	Sommittee y testing will only be performed on specimens round to be

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